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or stereoisomers or pharmaceutically acceptable salts, esters, or amides, wherein:

A is selected from NCH_2 , N(alkyl)CH_2 , CH_2N , $\text{CH}_2\text{N(alkyl)}$;

B is selected from H, $(\text{C}_3\text{-20})\text{alkyl}$, cycloalkyl, heteroalkyl, cycloalkylalkyl, heteroalkylalkyl, aryl, arylalkyl, heterocycle, heterocycloalkyl, each optionally substituted with R_1 and R_2 ;

D is selected from H, $(\text{C}_3\text{-20})\text{alkyl}$, cycloalkyl, heteroalkyl, cycloalkylalkyl, heteroalkylalkyl, aryl, arylalkyl, heterocycle, heterocycloalkyl, each optionally substituted with R_1 and R_2 ;

E is absent or selected from O, S, NH;

F is selected from N, NCH_2 , CH_2N ;

G is absent or selected from alkyl, alkyl interrupted by one or more heteroatoms, cycloalkyl, cycloalkyl interrupted by one or more heteroatoms;

J is absent or selected from aryl or heterocycle each optionally substituted with R_1 and R_2 ;

K is absent or selected from an alkyl, alkyl interrupted by one or more heteroatoms, cycloalkyl interrupted by one or more heteroatoms, cycloalkylalkyl interrupted by one or more heteroatoms, each optionally substituted with R_1 and R_2 ;

L is selected from H, chlorine, fluorine, bromine, iodine, OH, O(alkyl), amine, alkyl, fluoroalkyl, amide, NO_2 , SH, $\text{S(O)}_n(\text{alkyl})$, SO_3H , SO_3alkyl , aldehyde, ketone, acid, ester, urea, Oalkylamide, Oalkylester, Oalkylacid, Nalkylacid, alkylamine, alkylamide, alkylketone, alkylacid, alkylester, alkylurea, Nalkylamide,